Docket No. GJE-7705 Serial No. 10/591,137

## In the Claims

This listing of claims will replace all prior versions and listings of claims in this application.

1 (currently amended). A method for the treatment-or prevention of a condition associated with T-cell proliferation or that is mediated by pro- and/or anti-inflammatory cytokines, wherein said method comprises administering, to a patient in need of such treatment, a compound of formula (I)

wherein R<sub>1</sub> is H or Me;

 $R_2$  is H or alkyl and  $R_3$  is H or Me, or  $R_2$  and  $R_3$  are -CH<sub>2</sub>- thereby forming a ring; n is 0 to 2:

X is CH2 or O; and

the two benzene rings are each optionally substituted with OH, OMe, halogen, NHCHO, NHSO<sub>2</sub>Me, CONH<sub>2</sub>, SOMe, OCH<sub>2</sub>O or CH<sub>2</sub>OH;

and wherein the compound is selected from bufeniode, denopamine, fenoterol; ifenprodil, isoxsuprine, labetalol, medroxalol, mesuprine, nylidrin, protokylol, ractopamine, ritodrine, salmefamol and sulfinalol.

2 (currently amended). The method according to claim 1, wherein the condition is a ehronic degenerative disease such as rheumatoid arthritis, ostcoarthritis or ostcoporosis.

3 (withdrawn and currently amended). The method according to claim 1, wherein the condition is a chronic demyelinating disease such as multiple sclerosis. 4 (withdrawn and currently amended). The method according to claim 1, wherein the condition is a respiratory disease such as asthma or chronic obstructive pulmonary disease.

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- 5 (withdrawn and currently amended). The method according to claim 1, wherein the condition is an inflammatory bowel disease (IBD) such as ulcerative colitis or Crohn's disease.
- 6 (withdrawn and currently amended). The method according to claim 1, wherein the condition is a dermatological condition such as psoriasis, scleroderma or atopic dermatitis.
- 7 (withdrawn and currently amended). The method according to claim 1, wherein the condition is a dental disease such as periodontal disease or gingivitis.
- 8 (withdrawn). The method according to claim 1, wherein the condition is diabetic nephrophathy, lupus nephritis, IgA nephrophathy or glomerulonephritis.
- 9 (withdrawn and currently amended). The method according to claim 1, wherein the condition is systemic lupus erythematosus erythematosus (SLE).
- 10 (withdrawn). The method according to claim 1, wherein the condition is graft vs host disease.
  - 11 (withdrawn). The method according to claim 1, wherein the condition is a pain condition.
- 12 (withdrawn and currently amended). The method according to claim 11, wherein the pain condition is ehronic pain-such as chronic back pain, malignant pain, chronic headache (including migraine and cluster headaches) or arthritic pain).

13 (withdrawn and currently amended). The method according to claim 11, wherein the pain condition is acute pain such as post-operative pain, post-traumatic pain or acute disease-induced pain.

14 (withdrawn). The method according to claim 11, wherein the pain condition is neuropathic pain.

## 15 (cancelled).

16 (currently amended). The method according to claim 1, wherein the compound is in the form of the enantiomer or diastereomer that has relatively little or no activity at [[the]] an  $\alpha$  or  $\beta$  adrenoceptor.

## 17 (cancelled).

18 (new). A method for the treatment of a condition associated with T-cell proliferation or that is mediated by pro- and/or anti-inflammatory cytokines, wherein said method comprises administering, to a patient in need of such treatment, a compound of formula (I)

wherein R1 is H or Me;

 $R_2$  is H or alkyl and  $R_3$  is H or Me, or  $R_2$  and  $R_3$  are -CH2- thereby forming a ring; n is 0 to 2;

X is CH2 or O; and

the two benzene rings are each optionally substituted with OH, OMe, halogen, NHCHO, NHSO<sub>2</sub>Me, CONH<sub>3</sub>, SOMe, OCH<sub>3</sub>O or CH<sub>3</sub>OH;

wherein the compound is selected from bufeniode, denopamine, ifenprodil, isoxsuprine, labetalol, medroxalol, mesuprine, nylidrin, protokylol, ractopamine, ritodrine, salmefamol and sulfinalol:

wherein the compound is in the form of the enantiomer or diaster comer that has little or no activity at an  $\alpha$  or  $\beta$  adrenoceptor.